Attorney's Docket No.: 06275-0492US1 / 101170-1P

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Amendments to the Claims:

This listing of claims replaces all prior versions and listings of claims in the application:

Listing of Claims:

- 1. (Currently amended) A method of diagnosis comprising:
- (a) providing a biological sample from a human identified as being in need of treatment with a therapeutic agent that is transported by OATP-C rosuvastatin, wherein the sample comprises a nucleic acid encoding OATP-C;
 - (b) testing the nucleic acid for the presence, on at least one allele, of either
- (i) a codon encoding alanine at the position corresponding to position 174 of SEQ ID NO:1, or
 - (ii) an allele of a polymorphism in linkage disequilibrium with (i); and
- (c) if either (i) or (ii) is found in at least one allele, diagnosing the human as likely to have reduced ability to transport the therapeutic agent rosuvastatin into liver cells.
- 2. (Withdrawn) A method according to claim 1 wherein the polymorphism of (b)(ii) is 26A>G, -118A>C, -309T>C, -878A>G, -903C>T, -1054G>T, -1215T>A, or -1558T>C, all of SEQ ID NO:2; or T2122G, C2158T, A2525C, or G2651A, all of SEQ ID NO:3.
- 3. (Withdrawn) A method according to claim 1 wherein the polymorphism of (b)(ii) is selected from -118A>C and -1558T>C of SEQ ID NO:2.
- 4. (Currently amended) A method according to claim 1, wherein the therapeutic agent is a statin, the human is being treated with one dose level of <u>rosuva</u>statin and step (c) further comprises diagnosing the human as suitable for titration to another higher <u>rosuva</u>statin dose level

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comprising monitoring for a decrease in benefit risk ratio resulting from the reduced ability to transport the statin into cells if either (i) or (ii) is found in at least one allele.

5.-6. (Canceled)

- 7. (Currently amended) A method according to claim [[5]] 1 wherein the human is being treated with at least 5 mg of [[a]] rosuvastatin daily.
- 8. (Currently amended) A method according to claim [[5]] 1 wherein the human is being treated with at least 10 mg of [[a]] rosuvastatin daily.
- 9. (Currently amended) A method according to claim [[5]] 1 wherein the human is being treated with at least 20 mg of [[a]] rosuvastatin daily.
- 10. (Currently amended) A method according to claim [[5]] 1 wherein the human is being treated with at least 40 mg of [[a]] rosuvastatin daily.
- 11. (Currently amended) A method of diagnosis comprising:
- (a) providing a biological sample from a human identified as being in need of treatment with a therapeutic agent that is transported into cells by OATP-C rosuvastatin, wherein the sample comprises an OATP-C polypeptide;
- (b) determining whether the amino acid of the OATP-C polypeptide corresponding to position 174 of SEQ ID NO:1 is a valine; and
- (c) if the amino acid is not a valine, diagnosing the human as likely to have a reduced ability to transport the therapeutic agent rosuvastatin into liver cells.
- 12. (Currently amended) A method according to claim 11, wherein the therapeutic agent is a statin, the human is being treated with one dose level of <u>rosuva</u>statin and step (c) further

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comprises diagnosing the human as suitable for titration to another, higher <u>rosuva</u>statin dose level comprising monitoring for a decrease in benefit risk ratio resulting from the reduced ability to transport the statin into cells if the amino acid is not a valine.

- 13. (Withdrawn-Currently amended) A method according to claim [[12]] 11, the method further comprising measurement of measuring the level of OATP_C polypeptide expression with valine and/or alanine at position 174 whereby to determine the presence or absence of -118A>C polymorphism in OATP-C nucleic acid.
- 14. (Withdrawn-Currently amended) A method according to claim [[12]] 11, the method further comprising measuring OATP-C polypeptide for presence or absence of OATP-C *15 allele whereby to determine the presence or absence of -118A>C polymorphism in OATP-C nucleic acid determining, in a sample of nucleic acid from the human, the presence or absence, on at least one allele, of a cytosine at the position corresponding to -118 of SEQ ID NO:2, wherein the presence of the cytosine, combined with the determination that the amino acid of (b) is not a valine, is a further indication that the human is likely to have reduced ability to transport rosuvastatin into liver cells.
- 15. (Currently amended) A method according to claim [[12]] 11, wherein the amino acid at position 174 is determined to be alanine.
- 16. 17. (Canceled)
- 18. (Currently amended) A method according to claim [[16]] 11, wherein the human is being treated with at least 5 mg of [[a]] rosuvastatin daily.
- 19. (Currently amended) A method according to claim [[16]] 11, wherein the human is being treated with at least 10 mg of [[a]] rosuvastatin daily.

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20. (Currently amended) A method according to claim [[16]] 11, wherein the human is being treated with at least 20 mg of [[a]] rosuvastatin daily.

- 21. (Currently amended) A method according to claim [[16]] 11, wherein the human is being treated with at least 40 mg of [[a]] rosuvastatin daily.
- 22. (New) A method according to claim 1, wherein the nucleic acid is tested both for the presence, on at least one allele, of a codon encoding alanine at the position corresponding to position 174 of SEQ ID NO:1 and for the presence, on at least one allele, of a cytosine at the position corresponding to position -118 of SEQ ID NO:2.